

**EXHIBIT**

# Substance Sheet

Labjournal ID:

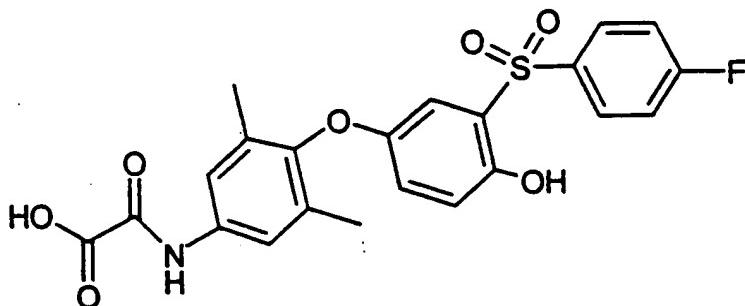
U-0282-85-B

Date:

Department: RES/MCD/MCDCHEM

Chem. Code:

US Sample No:

Chemists: KUKKOLA, PAIVI  
WANG, HUA

MW Eff.: 459.45

Factor f: 1.000

MF Subst.: C<sub>22</sub>H<sub>14</sub>FNO<sub>2</sub>S

MW Subst.: 459.45

Initial Amount: 50 mg

Subst./Stereo State:

Struct. Assign.:

Single cmpd - achiral  
Compatible with analytical  
data

Approval Status:

APPROVED

Synthesis Sheet:

Yes

End Product:

Yes

Multi.Par/CChem.:

No

To be sent to LFU/NCA: No

<sup>1</sup>H-NMR: Yes ; self service

MS: 2919-

IR: 2919-

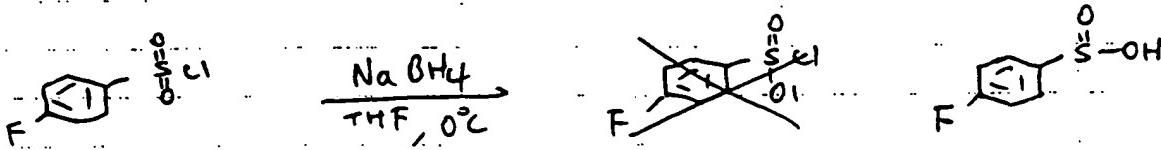
Solvent System: DMSO

Known in Lit.: No;

To be tested on: LPM; 4 mg, R. Steel

Clinical Codes:

Comment Batch: CHN; 100% purity by HPLC



	Amount	mmol	ef
4-fluorobenzensulfonyl chloride (194.61)	500 mg	2.57	1.0
Sodium borohydride (37.83)	0.49 g	12.85	5.0
THF	20 ml		

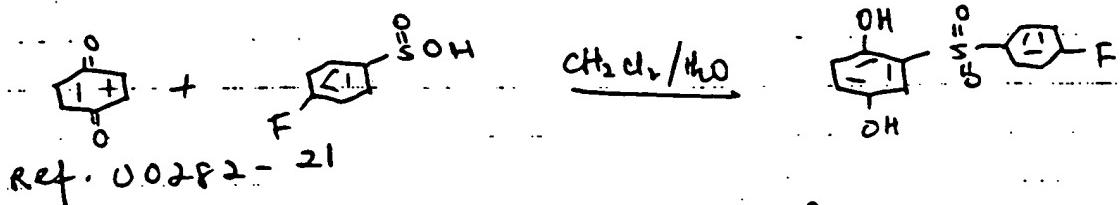
4-fluorobenzensulfonyl chloride was dissolved in THF and NaBH<sub>4</sub> was added in portions with stirring at 0°C. The rxn was stirred at 0°C for 1 h, then RT for 3 h.

hex/EtOAc	3:2
sm	sm

After removal of THF, H<sub>2</sub>O was added and mixture was acidified by the addition of HCl (6N) dropwise at 0°C. Extracted w/ EtOAc. Dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated to give 102.67 mg as a white solid.

MS: (m+) was detected

Unknown



U0282-67 (160)

$\text{CH}_2\text{Cl}_2$

1,4-benzoquinone (108.10)

H2O

Amount	mL	g
320 mg	2.0	1.05
6 mL		
206 mg	1.9	1.0
4 mL		

H2

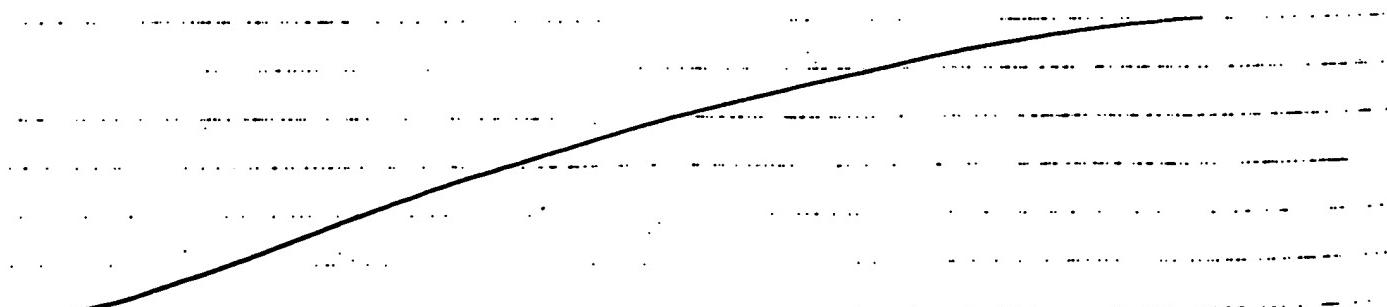
In a solution of U0282-67 in  $\text{CH}_2\text{Cl}_2$ , was added 1,4-benzoquinone

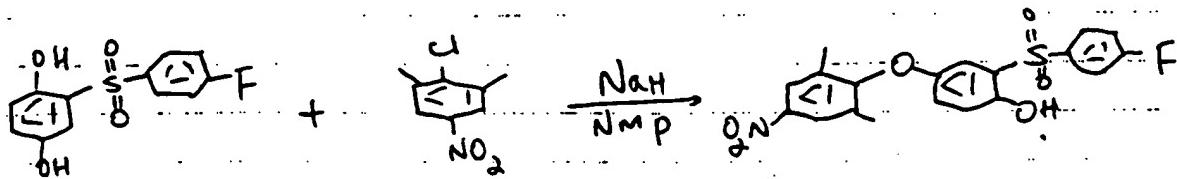
To a solution of 1,4-benzoquinone in  $\text{CH}_2\text{Cl}_2$ , was added a solution of U0282-67 in H2O. The solution was stirred at RT. After 10 min, precipitation ~~color~~ occurred. The suspension was stirred at RT for 3 h.

It was filtered. The solid was collected. #8 U0282-70 200mg as a off white solid

MS: AcN/H2O/NH4OH  
 $(2n-1)^-$  was detected

heating





Ref U022-25

U022-70 (268)

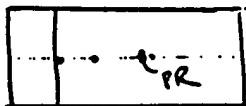
NaH (24, 607-1)

NB 2694-12 (185.6)

NMP

	Amount	mmol	of
200mg	0.75		1.0
99mg	2.46		3.3
167mg	0.90		1.2
5ml			

To a suspension of NaH in NMP at 0°C, was added U022-70, the suspension was stirred at RT for 30 min. NB 2694-12 was added. The black suspension was stirred at 0°C for 3h. It was quenched w/ PhO and extracted w/ Et<sub>2</sub>O ( $\times 3$ ). The organic layer was washed w/ brine, dried over Na<sub>2</sub>O<sub>4</sub>, and concentrated to give U022-74 as a yellow solid.

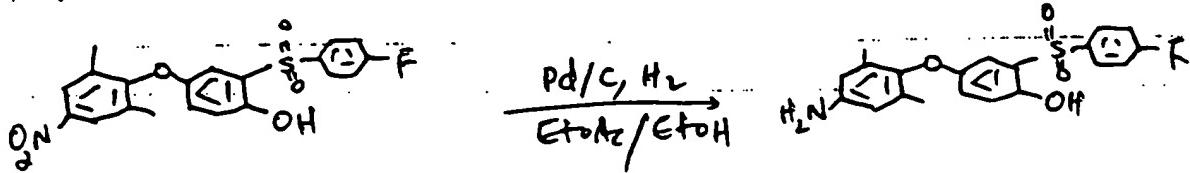
MS: (M-1)<sup>-</sup> was detected (416)

hex/EtOAc 3:2

The crude was purified w/ flash chro matography (hex/EtOAc 3:2) to give U022-74A. 246mg as a white solid.

<sup>1</sup>H-NMR (CDCl<sub>3</sub>) of U022-74A: reasonably clean

Hear



Ref. U0282-36

U0282-74A

Pd/C

EtOH/EtOAc

Ammonium mmol of  
240 mg  
24 mg  
conc / 10 ml

The rxn was stirred at RT f under th (crat) for 17 h.

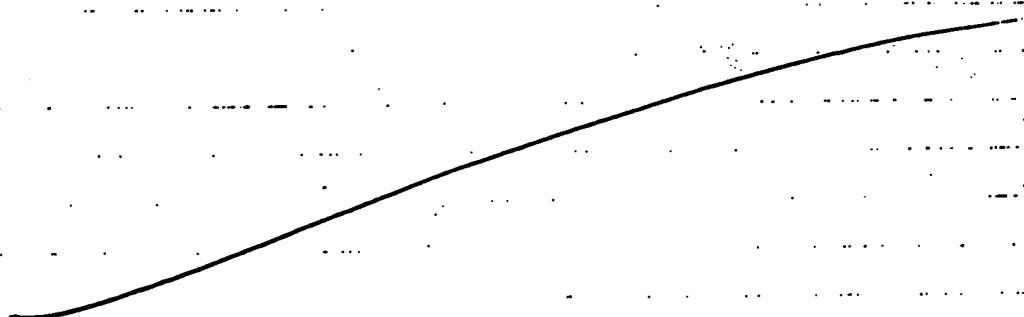
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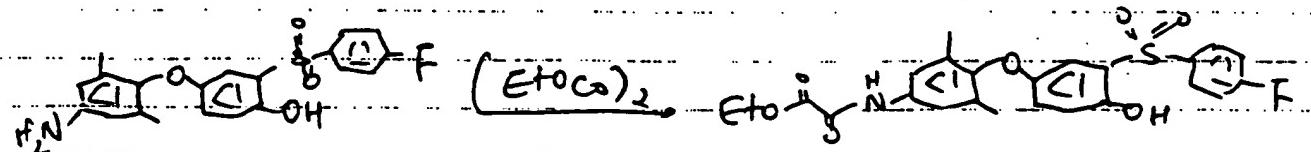
H2/EtOAc 2:3

Small amount of starting material still present. It was stirred under H<sub>2</sub> balloon at RT for 4 h. It was completed by 71%. It was filtered through celite. The filtrate was concentrated to give U0282-80 190 mg as a solid.

<sup>1</sup>H NMR (CDCl<sub>3</sub>): 0%

Area w -





Ref. U0282-38

U0282-80

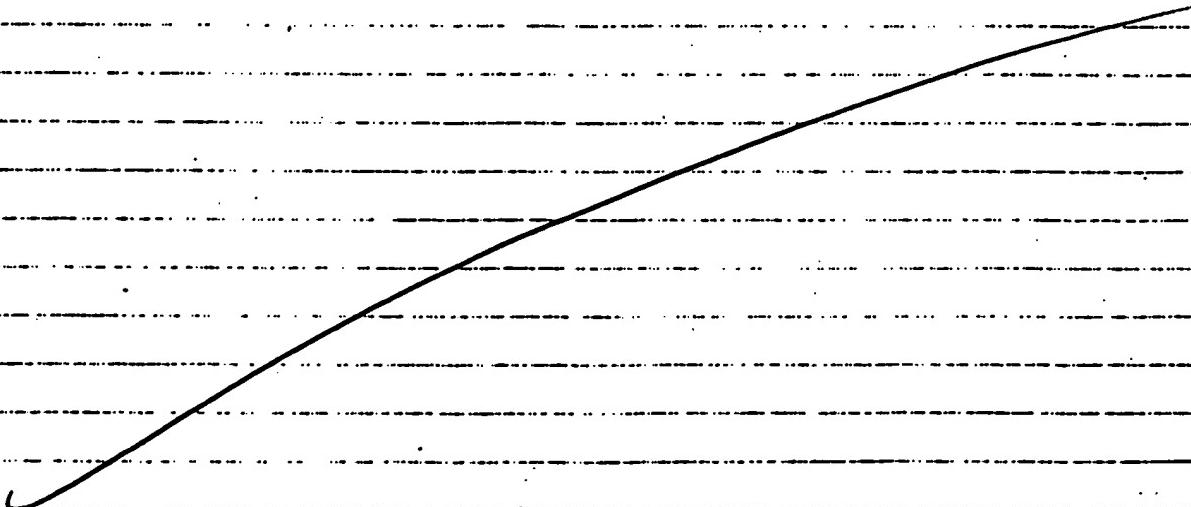
$(\text{EtOCo})_2$

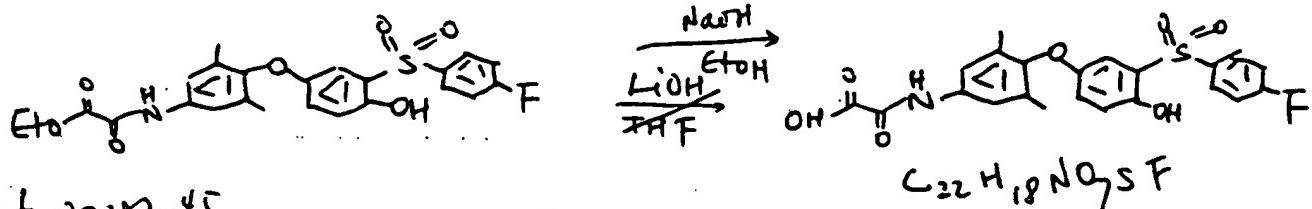
Amount mmol of  
1.90 mg  
2 ml.

The rxn was stirred at 180°C for 3 h. The solvents were removed by N<sub>2</sub> stream. Chromatographed on silica (hex/EtOAc 2:1) to give U0282-83A as a yellowish foam 195 mg.

<sup>1</sup>H NMR (CDCl<sub>3</sub>): OK, trace amount of impurities.

Fluor - ✓





Ref U02t2-45

U02t2-83 (487.5)

NaOH (1M)

EtOH

Amount	mmol	eq
170 mg	0.35	1.0
1.05 mL	1.05	3.0
3 mL		

The rxn was stirred at RT for 2 h.

10% MeOH in  $\text{CH}_2\text{Cl}_2$

It was quenched w/ HCl (1N, 1.5 mL). Extracted w/ EtOAc. The organ. was washed w/ brine. Dried over  $\text{Na}_2\text{SO}_4$  and concentrated to give U02t2-85 as a foam.

U02t2-85 was triturated w/  $\text{Et}_2\text{O}$ /hexane. Dried to give U02t2-85A. 75 mg as a white solid.

$^1\text{H-NMR}$  of U02t2-85A: same ether.

U02t2-85A was dried in vacuum over at 50°C for 1 h to give U02t2-85 57mg as a white solid.

$^1\text{H-NMR}$  ( $\text{CDCl}_3$ ): OK.

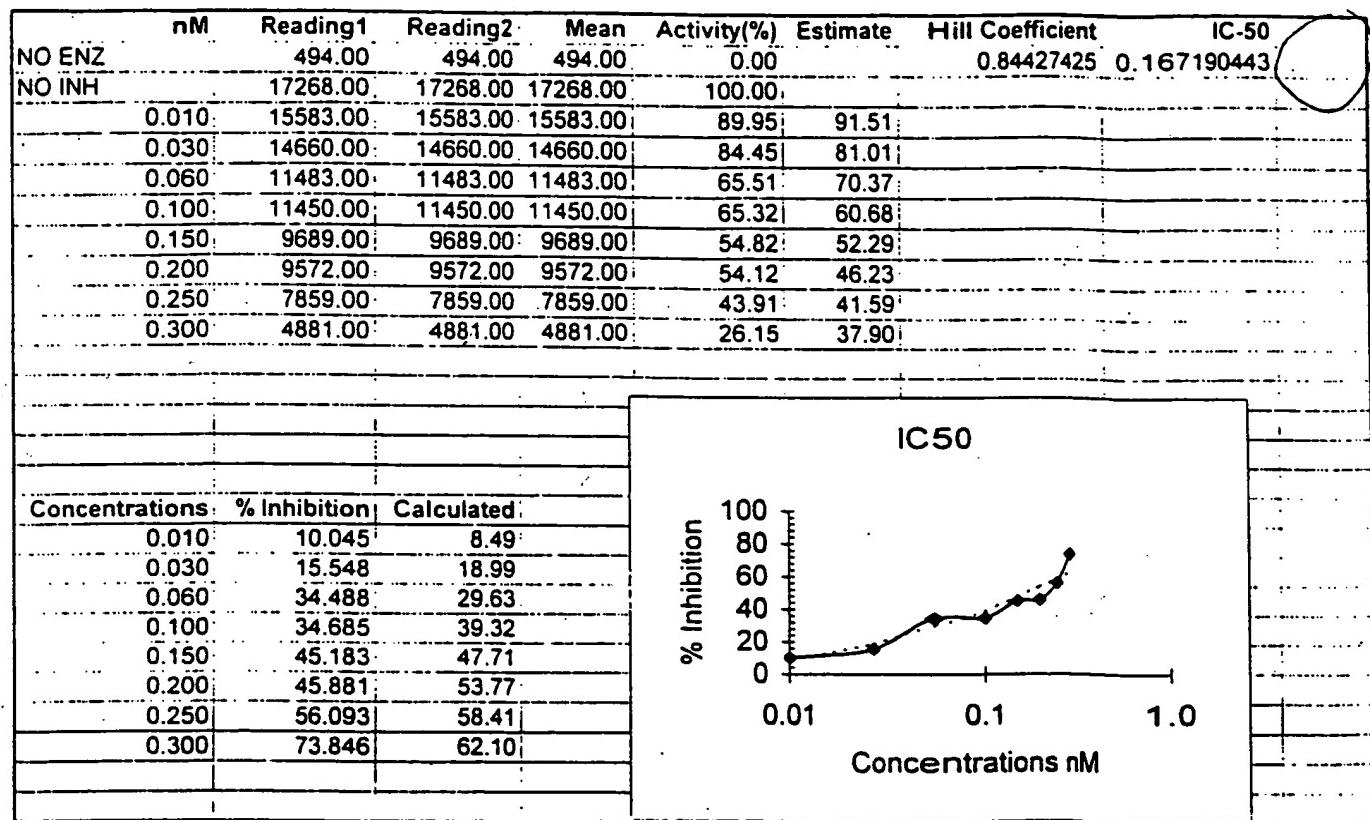
MS:  $(m-1)^{-}$  detected.

CHN: OK.

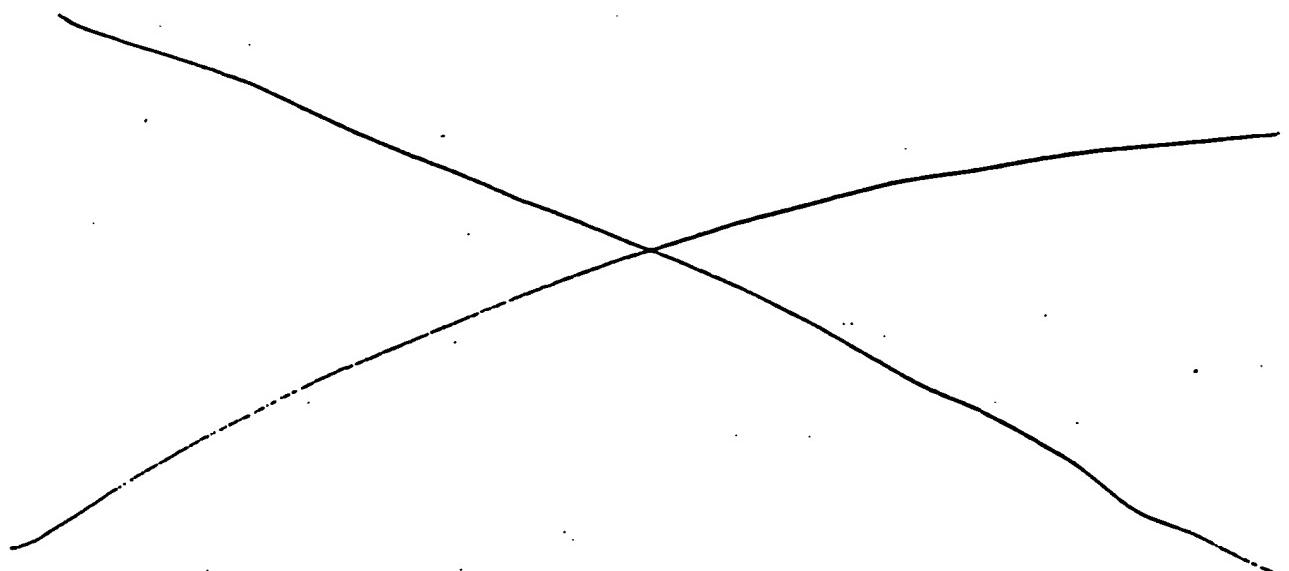
50mg sample sent out.

Hear

Sheet1



*Jorge J. Wenzel*



Read and understood by me

*Chidiebele*

Date